Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency’s estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology; and (e) estimates of capital or start-up costs and costs of operation, maintenance, and purchase of services to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology; and to review the collection of information; and to maintain and disclose, as required, the information. It does not mean the work already required of the public to produce the particular data. The voluntary nature of the collection is explained below.

Proposed Project

Data Collection for CDC Fellowship Programs—New—Division of Scientific Education and Professional Development (DSEPD), Centers for Surveillance, Epidemiology, and Laboratory Services (CSELS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC’s mission is to protect America from health, safety, and security threats, both foreign and in the U.S. To ensure a competent, sustainable, and empowered public health workforce...
CDC prepares to meet these challenges, CDC plays a key role in developing, implementing, and managing a number of fellowship programs. A fellowship is defined as a training or work experience lasting at least 1 month and consisting of primarily experiential (i.e., on-the-job) learning, in which the trainee has a designated mentor or supervisor. CDC fellowships are intended to develop public health professionals, enhance the public health workforce, and strengthen collaborations with partners in public health and healthcare organizations, academia, and other stakeholders in governmental and non-governmental organizations. Assessing fellowship activities is essential to ensure that the public health workforce is equipped to promote and protect the public’s health.

CDC requests a three-year approval of a generic clearance to collect data about its fellowship programs, as they relate to public health workforce development. Data collections will allow for ongoing, collaborative, and actionable communications between CDC fellowship programs and stakeholders (e.g., fellows, supervisors/mentors, alumni). These collections might include short surveys, interviews, and focus groups. Intended use of the resulting information is to:

- inform planning, implementation, and continuous quality improvement of fellowship activities and services;
- improve efficiencies in the delivery of fellowship activities and services; and
- determine to what extent fellowship activities and services are achieving established goals.

Collection and use of information about CDC fellowship activities will help ensure effective, efficient, and satisfying experiences among fellowship program participants and stakeholders.

CDC estimates that annually, a given fellowship program will conduct one query each with one of the three respondent groups: Fellowship applicants or fellows; mentors, supervisors, or employers; and alumni. The total annualized burden hours of 2,957 was determined as depicted in the following table.

### ESTIMATED ANNUALIZED BURDEN HOURS

<table>
<thead>
<tr>
<th>Type of respondent</th>
<th>Form name</th>
<th>Number of respondents</th>
<th>Total number of responses per respondent</th>
<th>Average burden per response (in hours)</th>
<th>Total burden (in hours)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Applicants or fellows</td>
<td>Fellowship Data Collection Instrument</td>
<td>1,848</td>
<td>1</td>
<td>30/60</td>
<td>924</td>
</tr>
<tr>
<td>Mentors, supervisors, or employers</td>
<td>Fellowship Data Collection Instrument</td>
<td>370</td>
<td>1</td>
<td>30/60</td>
<td>185</td>
</tr>
<tr>
<td>Alumni</td>
<td>Fellowship Data Collection Instrument</td>
<td>3,696</td>
<td>1</td>
<td>30/60</td>
<td>1,848</td>
</tr>
<tr>
<td>Total</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>2,957</td>
</tr>
</tbody>
</table>

LeRoy A. Richardson,
Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2016–14956 Filed 6–23–16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10458]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS’ intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish a notice in the Federal Register concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including any of the following subjects: the necessity and utility of the proposed information collection for the proper performance of the agency’s functions; the accuracy of the estimated burden; ways to enhance the quality, utility, and clarity of the information to be collected; and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments on the collection(s) of information must be received by the OMB desk officer by July 25, 2016.

ADDRESSES: When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395–5806 OR, Email: OIRA_submission@omb.eop.gov.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of the following:


2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov.

3. Call the Reports Clearance Office at (410) 786–1326.

FOR FURTHER INFORMATION CONTACT: Reports Clearance Office at (410) 786–1326.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term “collection of information” is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or